

FEB - 1 2001

K0/0091

 	SPECIAL 510(k) REAL PATIENT EEG WITH VIDEO
SPECIAL 510(k) NOTIFICATION OF A NEW DEVICE	JANUARY 16, 2001

510(k) Summary

This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of the SMDA 1990 and 21 CFR § 807.92.

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Director of R & D

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Common Names: Electroencephalograph

Classification:

Classification Name: Electroencephalograph
Product code: 84GWQ
Device Class: Class II
Regulation Number: 882.1400

Predicate Devices: Excel Tech EEG with Video [510(k) K000898]

Description: The Excel Tech Real Patient EEG with Video device is an EEG device with a parallel video stream.

The Real Patient EEG with Video allows for the distribution of EEG data via TCP/IP, Intranet, Internet, and physical means.

Substantial Equivalence:

The Excel Tech Real Patient EEG with Video is substantially equivalent to the Excel Tech EEG with Video [510(k) K000898]. The Real Patient EEG with Video allows for the distribution of EEG data via TCP/IP, Intranet, Internet, and physical means.

Indications for Use:

The Real Patient EEG with Video is intended to be used as an electroencephalograph: to acquire, display, store, archive electroencephalographic signals.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

FEB - 1 2001

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Ms. Sonja Markez
Regulatory Manager
Excel Tech, Ltd.
2568 Bristol Circle
Oakville, Ontario
Canada L6H 5S1

Re: K010091
Trade Name: EEG with Video
Regulatory Class: II
Product Code: GWQ
Dated: January 9, 2001
Received: January 11, 2001

Dear Ms. Markez:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the current Good Manufacturing Practice requirement, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic (QS) inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

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This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4595. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsmamain.html>".

Sincerely yours,

Miriam C. Provost
for Celia M. Witten, Ph.D., M.D.
Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure



SPECIAL 510(k) NOTIFICATION OF A NEW DEVICE

SPECIAL 510(k)
REAL PATIENT EEG
WITH VIDEO
JANUARY 16, 2001
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Statement of Indications for Use

Page 1 of 1

510(k) Number (if known): K010091

Device Name: EEG with Video

Indications for Use: The EEG with Video is an electroencephalograph which is designed to display the electrical signals of the brain on a video monitor.

(PLEASE DO NOT WRITE BELOW THIS LINE—CONTINUE ON ANOTHER
PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Miriam C. Purro
(Division Sign-Off)

Division of General Restorative Devices

510(k) Number K010091

Prescription Use ☒
(Per 21§ CFR 801.109)

OR Over-The Counter Use ☐

(Optional Format 1-2-96)